

Substitute for form 1449B/PTO
INFORMATION DISCLOSURE
STATEMENT BY APPLICANT

(use as many sheets as necessary)

Complete if Known

Application Number 10/522,130
Filing Date January 19, 2005
First Named Inventor Murray Goodman
Group Art Unit
Examiner Name
Attorney Docket Number SDUC1100J-1 (041673-3109)

Sheet 1 of 2

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No. ¹	U.S. Patent Document		Name of Patentee or Applicant of Cited Document	Date of Publication of Cited Document MM-DD-YYYY	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
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FOREIGN PATENT DOCUMENTS

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		Office ³	Number ⁴	Kind Code ⁵ (if known)				
	A1	WO	91/09958		Whitehead Institute for Biomedical Research	07-11-1991		
	A2	WO	94/04686		Biogen, Inc.	03-03-1994		
	A3	WO	98/52614		The Board of Trustees of the Leland Stanford Junior University	11-26-1998		

NON PATENT LITERATURE DOCUMENTS

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	A4	BUSCHLE, M. et al., Transloading of tumor antigen-derived peptides into antigen-presenting cells. Proc. Natl. Acad. Sci. USA., 94, p. 3256-3261 (1997).	
	A5	EMI, N. et al., Gene Transfer Mediated by Polyarginine Requires a Formation of Big Carrier-Complex of DNA Aggregate, Biophys. Res. Commun., 231, p. 421-424 (1997).	
	A6	FEICHTINGER, L. et al., Triurethane-Protected Guanidines and Triflydiurethane-Protected Guanidines: New Reagents for Guanidinylation Reactions, J. Org. Chem., 63, p. 8432 (1998).	
	A7	LEONETTI, J. -P. et al., Biological Activity of Oligonucleotide-Poly(L-lysine) Conjugates: Mechanism of Cell Uptake, Bioconjugate Chem., 1, p. 149-153 (1990).	
	A8	MITCHELL, D.J. et al., Polyargine enters cells more efficiently than other polycationic homopolymers, J. Peptide Res., 55 p. 318-325 (2000).	
	A9	MURPHY, J.E. et al., A combinatorial approach to the discovery of efficient cationic peptoid reagents for gene delivery, Proc. Natl. Acad. Sci. USA., 95, p. 1517-1522 (1998).	

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*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609. Draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.

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	A10	PEPINSKY, R.B. et al., Specific Inhibition of a Human Papillomavirus E2 <i>Trans</i> -Activator by Intracellular Delivery of Its Repressor, DNA Cell Biol., 13, p. 1011-1019 (1994).	
	A11	RYSER, H.J.-P., A Membrane Effect of Basic Polymers dependent on Molecular Size, Nature (London), 215, p. 934-936 (1967).	
	A12	RYSER, N.J. -P. et al., Conjugation of methotrexate to poly(L-lysine) increases drug transport and overcomes drug resistance in cultured cells, Proc. Nat. Acad. Sci. USA., 75, p. 3867-3870 (1978).	
	A13	SCHWARZE, S.R. et al., In Vivo Protein Transduction: Delivery of a Biologically Active Protein into the Mouse, Science, 285, p. 1569-1572 (1999).	
	A14	SHEN, W., et al., Conjugation of poly-L-lysine to albumin and horseradish peroxidase: A novel method of enhancing the cellular uptake of proteins, Proc. Nat. Acad. Sci. USA., 75, p. 1872-1876 (1978).	
	A15	VOCERO-AKBANI, A.M. et al., Killing HIV-infected cells by transduction with an HIV protease-activated caspase-3 protein, Nat. Med., 5, p. 29-33 (1999).	
	A16	WENDER, P.A. et al., The design, synthesis, and evaluation of molecules that enable or enhance cellular uptake: Peptoid molecular transporters, Proc. Natl. Acad. Sci. USA., 97, p. 13003-13008 (2000).	

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